

**Amendments to the Specification:**

Please amend the specification as follows:

Please replace paragraph starting at page 1, lines 26-32, with the following rewritten paragraph:

These are major problems in the industrial development of human or humanized chimeric monoclonal antibodies. By way of example, the company Protein Design Labs has suspended phase I/II clinical trials of ~~Remitogen®~~ REMITOGEN® (apolizumab), which is an anti-HLA-DR antibody that may be used for treating cancers of MHC class II-positive cells, in particular B-cell and T-cell leukemias.

Please replace paragraph starting at page 3, line 29 to page 4, line 1, with the following rewritten paragraph:

The invention therefore proposes antibodies which have an activity that is up to 100 times greater than the available antibodies in therapy. In particular, the invention provides an anti-HLD-DR and an anti-CD20 that are significantly more effective than their respective homologue such as ~~Remitogen®~~ REMITOGEN® (apolizumab) and ~~Rutixan®~~ RITUXAN® (Rituximab). The present invention marks a major turning point in the development of antibodies for clinical purposes, by providing a new generation, the effective doses of which in order to obtain 50% activity are much lower than those of the antibodies currently used.

Please replace Table 1 starting at page 8, line 21, and ending at page 10, with the following rewritten Table 1:

Antibody name and trade mark	Company	Target	Indication
<del>Edrecolomab</del> <u>PANOREX</u> (Edrecolomab)	Centocor	anti-Ep-CAM	colorectal cancer
<del>Rituximab</del> <u>RITUXAN</u> (Rituximab)	Idec licensed to Genentech/ Hoffmann la Roche	anti-CD20	B cell lymphoma thrombocytopenia purpura

<del>Trastuzumab</del> <u>HERCEPTIN</u> <del>Trastuzumab</del>	Genentech licensed to Hoffmann la Roche/ Immunogen	anti-HER2	ovarian cancer
<del>Palivizumab</del> <u>SYNAGIS</u> <del>(Palivizumab)</del>	Medimmune licensed to Abott		RSV
<del>Alemtuzumab</del> <u>CAMPATH</u> <del>(Alemtuzumab)</del>	BTG licensed to Schering	anti-CD52	leukemia
<u>ZEVALIN</u> <del>(ibritumomab tiuxetan)</del>	IDEC licensed to Schering	anti-CD20	NHL
<del>Cetuximab</del> <u>IMC-C225</u> <del>(Cetuximab)</del>	Merck/BMS/ Imclone	anti-HER1	cancers
<del>Bevacizumab</del> <u>AVASTIN</u> <del>(Bevacizumab)</del>	Genentech/ Hoffmann la Roche	anti-VEGFR	cancers
Epratuzumab	Immumedics/ Amgen	anti-CD22	cancers: non-Hodgkin lymphoma
Hu M195Mab	Protein Design Labs	ND	cancers
MDX-210	Immuno-Designed Molecules	ND	cancers
BEC2 <del>(Mitumomab)</del>	Imclone	anti-GD3	cancers
<del>Oregovomab</del> <u>OVAREX</u> <del>(Oregovomab)</del>	Altarex	anti-CA125	Ovarian cancer
<del>Ecrimeximab</del> KW-2971 <del>(Ecrimeximab)</del>	Kyowa-Hakko	anti-GD	malignant melanoma
ABX-EGF	Abgenix	EGF	cancers
MDX010	Medarex	ND	cancers
XTL 002	XTL bio-pharmaceuticals	ND	anti-viral: HCV
H11 SCFV	viventia biotech	ND	cancers
4B5	viventia biotech	anti-GD2	cancers
XTL 001	XTL biopharmaceuticals	ND	anti-viral: HBV

MDX-070	MEDAREX	Anti-PSMA	Prostate cancer
TNX-901	TANOX	anti-CD23	
IDEC-114	IDEC	inhibition ProteinC	non-Hodgkin lymphoma

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